

**UNITED STATES DISTRICT COURT
FOR THE SOUTHERN DISTRICT OF NEW YORK**

DORIS KAMINSKY, as Executor of the Estate of
ROBERT KAMINSKY, and DORIS KAMINSKY,
individually,

Plaintiff,

v.

KONINKLIJKE PHILIPS N.V.; PHILIPS
NORTH AMERICA LLC; PHILIPS HOLDING
USA, INC.; and PHILIPS RS NORTH AMERICA
LLC,

Defendants.

**COMPLAINT AND DEMAND
FOR JURY TRIAL**

Civil Case No. _____

CIVIL COMPLAINT

Plaintiff, Doris Kaminsky, as Executor of the Estate of Robert Kaminsky, and Doris Kaminsky, individually, residing in Ulster County in the State of New York, by and through Plaintiff's attorneys, Merson Law, PLLC, hereby submits the following Complaint and Demand for Jury Trial against Defendants Koninklijke Philips N.V. ("Royal Philips"), Philips North America LLC ("Philips NA"), Philips Holding USA, Inc. ("Philips USA") and Philips RS North America LLC ("Philips RS") (hereinafter, collectively referred to as "Philips" or "Defendants") and alleges upon information and belief that at all relevant times hereinafter mentioned:

NATURE OF ACTION

1. Philips manufactures, markets, sells, and distributes a variety of products for sleep and home respiratory care.

2. Philips manufactures, markets, imports, sells, and distributes a variety of Continuous Positive Airway Pressure (CPAP) and BiLevel Positive Airway Pressure (BiPAP) devices for patients with sleep apnea.

3. Philips manufactures, markets, imports, sells, and distributes a variety of ventilator devices for patients with respiratory conditions.

4. On April 26, 2021, as part of its Quarterly Report for Q1 2021, under a section entitled “Regulatory Update,” Philips publicly disclosed (for the first time) that the sound abatement foam in Philips’ CPAP, BiPAP, and mechanical ventilator devices posed serious health risks to their users.

5. On June 14, 2021, Philips issued a recall notification for (many of) its CPAP, BiPAP, and mechanical ventilator devices.

6. In its recall notification, Philips advised of potential health risks related to the sound abatement foam used in the affected devices.

7. Philips’ recall advised that patients using these (affected) devices of potential risks from exposure to chemicals released from the sound abatement foam via degradation and/or off-gassing.

8. Specifically, Philips’ recall notification stated that the risks related to exposure to chemicals given off/by the sound abatement foam could include headache, irritation, inflammation, respiratory issues, and possible toxic and carcinogenic effects.

9. Decedent Robert Kaminsky (hereinafter “Decedent”) (first) was prescribed the use of and purchased Philips’ devices, the DreamStation CPAP machine and the Philips System One, to treat his sleep apnea in 2012.

10. Decedent used Philips’ DreamStation CPAP machine and the Philips System One devices (hereinafter, the “Subject Devices” or “Devices”) essentially on a daily basis for a number

of years.

11. As a direct and proximate result of Philips' Devices, Decedent suffered serious and substantial life-altering injuries including the development of T-cell/histiocyte-rich large B-cell lymphoma.

12. As a direct and proximate result of the Subject Devices manufactured, marketed, sold, and distributed by Philips, Decedent suffered physical and emotional injuries, including cancer and the invasive treatment of the cancer and the sequelae of the cancer and its treatment.

13. As a direct and proximate result of Decedent's use of Philips' Devices, Decedent died on February 18, 2023.

14. As a direct and proximate result of Decedent's use of Philips' Devices, plaintiff Doris Kaminsky was deprived of the services, society and companionship of her husband (Decedent) and was caused to become obliged to expend sums of money for medical and hospital care on his behalf.

15. Defendants have long known that the polyester-based polyurethane (PE-PUR) sound-abatement foam in Defendants' CPAP, BiPAP, and mechanical ventilator devices (have tendencies to) release toxic and carcinogenic microparticles that can be inhaled by users like Decedent, causing serious injury or death.

16. As a result of the Devices' defects and Defendants' tortious acts/omissions, Decedent developed a serious and life-threatening cancer, endured unnecessary pain and suffering as well as lost wages and a diminished capacity to earn wages, and subsequently died.

17. Decedent suffered from unnecessary pain, debilitation, hospitalization, and the development of Hodgkin's lymphoma and death, and Plaintiff Doris Kaminsky lost her husband because Defendants defectively designed the Devices and failed to adequately warn of the dangers

of the Devices.

PARTIES

18. At all relevant times, Plaintiff Doris Kaminsky is a resident and citizen of Ulster County in the State of New York. Decedent Robert Kaminsky was injured and subsequently caused his death due to the defective medical Devices manufactured by Defendants.

19. Plaintiff Doris Kaminsky was duly appointed executor of the Estate of Robert Kaminsky by Order of the Surrogate's County of the State of New York, Ulster County dated April 21, 2023.

20. Defendant Koninklijke Philips N.V. ("Royal Philips") is a public limited liability company established under the laws of The Netherlands, having its principal executive offices at Philips Center, Amstelplein 2, 1096 BC Amsterdam, The Netherlands. Royal Philips is the parent company of Philips NA and Philips RS. Royal Philips can be served with process via the *Convention on the Service Abroad of Judicial and Extrajudicial Documents in Civil or Commercial Matters* ("the Hague Convention").

21. Defendant Philips North America LLC ("Philips NA") is a Delaware corporation with its principal place of business located at 222 Jacobs Street, Floor 3, Cambridge, Massachusetts 02141. Philips NA is a wholly-owned subsidiary of Royal Philips. Upon information and belief, Philips NA manages the operation of Royal Philips' various lines of business, including Philips RS, in North America. The sole member of Philips NA is Philips USA.

22. Defendant Philips Holding USA, Inc. ("Philips USA") is a Delaware corporation with its principal place of business located at 222 Jacobs Street, Floor 3, Cambridge, Massachusetts 02141. Philips USA is a holding company that is the sole member of Defendant Philips NA.

23. Defendant Philips RS North America LLC (“Philips RS”) is a Delaware corporation with its principal place of business located at 6501 Living Place, Pittsburgh, Pennsylvania 15206. Philips RS was formerly operated under the business name Respiroics, Inc. (“Respiroics”). Royal Philips acquired Respiroics in 2008.

STATEMENT OF JURISDICTION & VENUE

24. This Court has subject matter jurisdiction over the parties pursuant to 28 U.S.C. § 1332(a) because plaintiff and defendants are citizens of different states and the amount in controversy exceeds \$75,000.00, exclusive of interest and costs.

25. Venue is proper in this Court pursuant to 28 U.S.C. § 1391(a) and (c).

26. Upon information and belief, at all relevant times, Defendants were present and regularly transacted, solicited, and conducted business in the Southern District of New York, through its employees, agents, and/or sales representatives, purposefully availing themselves of the privilege of doing business in New York and deriving substantial revenue from such business.

27. At all relevant times, Defendants placed the defective Devices into the stream of interstate commerce that were purchased and used by Decedent.

28. Defendants are conclusively presumed to have been doing business in this state and are subject to New York’s long arm jurisdiction.

29. At all relevant times, Defendants expected or should have expected that their acts and omissions would have consequences within the United States and Southern District of New York.

30. This Court has personal jurisdiction over Defendants because of their systematic and continuous contacts with New York and such exercise of personal jurisdiction over Defendants comports with due process.

FACTUAL BACKGROUND

31. At all relevant times, Defendants manufactured, sold, and distributed a line of CPAP and BiPAP devices as well as mechanical ventilator devices under its “Sleep & Respiratory Care” portfolio. These devices are designed to assist individuals with a number of sleep, breathing, and respiratory conditions, including sleep apnea.

32. Defendants sought and obtained clearance from the Food and Drug Administration (“FDA”) to market the recalled devices, including the Devices used by Plaintiff, under Section 510(k) of the Medical Device Amendment to the Food, Drug, and Cosmetics Act (“FDCA”). Section 510(k) allows marketing of medical devices if the device is deemed substantially equivalent to other legally marketed predicated devices. Obtaining clearance under 510(k) is significantly less rigorous than through the pre-market approval (“PMA”) process, as no formal review for safety or efficacy is performed and no clinical data is required.

A. Continuous Positive Airway Pressure (CPAP) Therapy

33. Continuous Positive Airway Pressure (“CPAP”) therapy is a common nonsurgical treatment primarily used to treat sleep apnea. CPAP therapy typically involves the use of a nasal or facemask device and a CPAP machine, which helps individuals breathe by increasing the air pressure in an individual’s throat.

34. Sleep apnea is a common sleep disorder affecting millions of Americans, including Plaintiff, and characterized by repeated interruptions in breathing through an individual’s sleep cycle. These interruptions, called “apneas,” are caused when the soft tissue in an individual’s airway collapses. The airway collapse prevents oxygen from reaching the individual’s lungs which can cause a buildup of carbon dioxide. When the individual’s brain senses the buildup of carbon dioxide, it briefly rouse the individual from sleep so that the individual’s airway can reopen. Often these interruptions are so brief that the individual will not remember. Despite the brevity of the

interruption, the sleep cycle disruption caused by sleep apnea can dramatically impact a person's lifestyle, including negative impacts to energy levels, mental performance, and long-term health. CPAP therapy helps treat sleep apnea by forcing pressurized air through the individual's airway, preventing the individual's airway from collapsing while breathing during sleep cycles, which can help prevent interruptions in breathing.

B. Bi-Level Positive Airway Pressure (BiPAP) Therapy

35. Bi-Level Positive Airway Pressure ("BiPAP") therapy is a common alternative to CPAP therapy for treating sleep apnea. Similar to CPAP therapy, BiPAP therapy is nonsurgical and involves the use of a nasal or facemask device to maintain air pressure in an individual's airway. BiPAP is distinguishable from CPAP therapy, in that BiPAP devices deliver two alternating levels—inspiratory and expiratory—of pressurized air into a person's airway, rather than the single continuous level of pressurized air delivered by a CPAP device. The inspiratory positive airway pressure assists a person as a breath is taken in. Conversely, the expiratory positive airway pressure is applied to allow a person to comfortably breathe out. BiPAP devices deliver one level of pressurized air (the inspiratory positive level) to assist as a person inhales, and another level (the expiratory level) as a person exhales.

C. Philips' Sleep & Respiratory Care Devices Were Endangering Users

36. On April 26, 2021, as part of its Quarterly Report for Q1 2021, Philips publicly disclosed (for the first time), under a section titled "Regulatory Update," that device user reports had led to a discovery that the type of PE-PUR "sound abatement" foam Philips used to minimize noise in several CPAP, BiPAP, and mechanical ventilator devices posed health risks to their users. Specifically, Philips disclosed that "the [PE-PUR] foam may degrade under certain circumstances, influenced by factors including the use of unapproved cleaning methods, such as ozone[], and

certain environmental conditions involving high humidity and temperature.”

37. Philips utilized polyester-based polyurethane (PE-PUR) sound abatement foam to dampen device vibration and sound during routine operation.

38. On June 14, 2021, almost two months after Philips notified its stockholders, it finally advised the medical community, medical equipment suppliers and some patients, by issuing a recall notification of specific devices allegedly based upon extensive ongoing review following the announcement on April 26, 2021.

39. In its recall notification, Philips identified examples of potential risks which include exposure to chemicals emitted from the sound abatement foam material via degradation and/or off-gassing.

40. Philips reported that, based on lab testing and evaluations, it may be possible that these (potential) health risks could result in a wide range of (potential) patient impact, from transient

potential injuries, symptoms, and complications, as well as possibly serious injury, which can be life-threatening or cause permanent impairment, or require medical intervention to preclude permanent impairment.

41. According to Philips’ recall notice, the PE-PUR foam used in recalled devices such as the Devices used by Plaintiff puts users at risk of suffering from the following health harms: “headache, irritation [skin, eye, and respiratory tract], inflammation, respiratory issues, and possible toxic and carcinogenic effects.”

42. On June 14, 2021, Philips also issued a brief report titled “Clinical Information for Physicians.” In that report, Philips disclosed that “[l]ab analysis of the degraded foam reveals the presence of potentially harmful chemicals including:

-Toluene Diamine

-Toluene Diisocyanate

-Diethylene glycol.”

43. In its report titled “Clinical Information for Physicians,” Philips also disclosed that lab testing performed by and for Philips had also identified the presence of Volatile Organic Compounds (VOCs) which may be emitted from the sound abatement foam component of the affected devices, stating “VOCs are emitted as gases from the foam included in the [affected devices] and may have short- and long-term adverse health effects. Standard testing identified two compounds of concern may be emitted from the foam that are outside of safety thresholds. The compounds identified are the following:

-Dimethyl Diazine

-Phenol, 2,6-bis (1,1-dimethylethyl)-4-(1-methylpropyl)-

D. Philips’ Recalled Devices

44. In total, Philips announced that “[b]etween 3 million and 4 million” devices are targeted in the recall.

45. The list of devices recalled by Philips (the “Recalled Devices”), which include the devices used by Decedent, include:

Philips CPAP and BiPAP Devices Subject to Recall	
Device Name/Model	Type
Philips E30 (Emergency Use Authorization)	Continuous Ventilator, Minimum Ventilatory Support, Facility Use
Philips DreamStation ASV	Continuous Ventilator, Non-life Supporting
Philips DreamStation ST, AVAPS	Continuous Ventilator, Non-life Supporting
Philips SystemOne ASV4	Continuous Ventilator, Non-life Supporting
Philips C Series ASV, S/T, AVAPS	Continuous Ventilator, Non-life Supporting
Philips OmniLab Advanced Plus, In-Lab Titration Device	Continuous Ventilator, Non-life Supporting

Philips SystemOne (Q Series)	Non-continuous Ventilator
Philips DreamStation, CPAP, Auto CPAP, BiPAP	Non-continuous Ventilator
Philips DreamStation GO, CPAP, APAP	Non-continuous Ventilator
Philips Dorma 400, 500, CPAP	Non-continuous Ventilator
Philips REMStar SE Auto, CPAP	Non-continuous Ventilator

Philips Device Name/Model	Type
Philips Trilogy 100 Ventilator	Continuous Ventilator
Philips Trilogy 200 Ventilator	Continuous Ventilator
Philips Garbin Plus, Aeris, LifeVent Ventilator	Continuous Ventilator
Philips A-Series BiPAP Hybrid A30	Continuous Ventilator, Minimum Ventilatory Support, Facility Use
Philips A-Series BiPAP V30 Auto Ventilator	Continuous Ventilator, Minimum Ventilatory Support, Facility Use
Philips A-Series BiPAP A40	Continuous Ventilator, Non-life Supporting
Philips A-Series BiPAP A30	Continuous Ventilator, Non-life Supporting

46. Philips issued the following advice to patients using any of the recalled devices:

- “For patients using BiLevel PAP and CPAP devices: Discontinue use of affected units and consult with physicians to determine the benefits of continuing therapy and potential risks.”
- “For patients using life-sustaining mechanical ventilator devices: DO NOT discontinue or alter prescribed therapy, without consulting physicians to determine appropriate next steps.”

E. Philips Unreasonably Delayed the Recall

47. Defendants have not disclosed when they first received reports from users of its Sleep & Respiratory Care devices “regarding the presence of black debris/particles within the airpath circuit (extending from the device outlet, humidifier, tubing, and mask).” However, given how long ago the first of the recalled devices came to market, (it is likely) upon information and belief Defendant learned of these issues for a substantial period of time before the recall.

Additionally, Philips released its next-generation CPAP device, the DreamStation 2, which does not have the defective, carcinogenic foam, on April 13, 2016 – not even two full weeks before Philips first publicly disclosed in its Q1 2021 Quarterly Report a potential health issue with its CPAP devices, including Plaintiff's defective first-generation DreamStation device. Defendants first sought FDA clearance for the DreamStation 2 in February 2020, and in all likelihood began developing it long before then.

48. Thus, as a result of user reports and other testing performed by and on behalf of Defendants, Defendants were aware of the degradation and off-gassing of the PE-PUR sound abatement foam used in the recalled devices, including Plaintiff's Device, yet continued to manufacture, market, and sell the recalled devices with such cognizance for a significant period of time. During this period, Defendants unreasonably and unjustly profited from the manufacture and sale of the recalled devices and unreasonably put users of the recalled devices at risk of developing adverse health effects, including cancer.

49. Decedent Robert Kaminsky was prescribed and purchased a Philips One System CPAP device in March 2012.

50. Decedent Robert Kaminsky was prescribed and purchased a Philips DreamStation CPAP device in March 2016.

DECEDENT ROBERT KAMINSKY'S DREAMSTATION CPAP DEVICE

51. Plaintiff Doris Kaminsky brings this product liability personal injury and wrongful death action as a recipient of defective medical devices, i.e., a CPAP device designed, manufactured and distributed by Defendants.

52. In March 2012, Decedent Robert Kaminsky was prescribed and purchased a Philips System One CPAP device.

53. In March 2016, Decedent Robert Kaminsky was prescribed and purchased a Philips DreamStation CPAP device.

54. Defendants, directly or through their subsidiaries or affiliates, designed, manufactured, distributed and sold the Philips CPAP devices prescribed to and purchased by Decedent Robert Kaminsky.

55. Based upon the patient population that Defendants intended its Philips System One and DreamStation CPAP devices to be used by, when Decedent Robert Kaminsky used the Devices, he was an appropriate patient to use the Devices.

56. At all times subsequent to the date of first use, Decedent Robert Kaminsky used the Devices in a normal and expected manner.

57. Subsequent to Plaintiff's normal use of the Devices, Decedent suffered symptoms including but not limited to increasing pain, discomfort, and the development of T-cell/histiocyte-rich large B-cell lymphoma which caused his death on February 18, 2023.

58. At the time the Devices were purchased by Decedent Robert Kaminsky, it was in the same condition in all relevant respects as when it left Philips' control.

59. Prior to Decedent Robert Kaminsky's purchase of the Devices in March 2012 and March 2016, Philips did not warn patients, including Decedent Robert Kaminsky, physicians, its customers, or its sales representative/distributors that the Devices were known to emit toxic and/or carcinogenic particles from its PE-PUR sound abatement foam via degradation and/or off-gassing, which could then be directly inhaled by the user, causing severe injury or death.

60. Decedent's use of the Devices subjected him to much greater risks of harm of personal injury and death than he had before using the Devices.

61. Had Decedent Robert Kaminsky or Plaintiff's physician known that the Devices would release carcinogenic particles causing Decedent's development of lymphoma, and death

then neither Decedent nor Decedent's physician would have chosen the Devices for treatment of Decedent's sleep apnea.

62. As a direct and proximate result of use of Philips' CPAP devices, Decedent Robert Kaminsky suffered significant harm, including but not limited to:

- (a) the development of fatal T-cell/histiocyte-rich large B-cell lymphoma and its invasive treatment;
- (b) pain and anguish, both in mind and in body;
- (c) permanent diminishment of his ability to participate in and enjoy the affairs of life;
- (d) medical bills associated with the treatment of Hodgkin's lymphoma and recovery therefrom;
- (e) future medical expenses;
- (f) loss of enjoyment of life;
- (g) loss of past and future earnings and earning capacity;
- (h) disfigurement;
- (i) physical impairment;
- (j) death.

FEDERAL STATUTORY AND REGULATORY REQUIREMENTS

63. Pursuant to federal law, a medical device is deemed to be adulterated if, among other things, it fails to meet established performance standards, or if the methods, facilities or controls used for its manufacture, packing, storage or installation are not in conformity with federal requirements. 21 U.S.C. § 351.

64. Pursuant to federal law, a device is deemed to be misbranded if, among other things, its labeling is false or misleading in any particular, or if it is dangerous to health when used in the

manner prescribed, recommended or suggested in the labeling thereof. 21 U.S.C. § 352.

65. Pursuant to federal law, manufacturers are required to comply with FDA regulation of medical devices, including FDA requirements for records and reports, in order to prohibit introduction of medical devices that are adulterated or misbranded, and to assure the safety and effectiveness of medical devices. In particular, manufacturers must keep records and make reports if any medical device may have caused or contributed to death or serious injury, or if the device has malfunctioned in a manner likely to cause or contribute to death or serious injury. Federal law also mandates that the FDA establish regulations requiring a manufacturer of a medical device to report promptly to FDA any correction or removal of a device undertaken to reduce a risk to health posed by the device, or to remedy a violation of federal law by which a device may present a risk to health. 21 U.S.C. § 360(i).

66. Pursuant to federal law, the Secretary of Health and Human Services may prescribe regulations requiring that the methods used in, and the facilities and controls used for, the manufacture, pre-production design validation (including a process to assess the performance of a device, but not including an evaluation of the safety or effectiveness of a device), packaging, storage and installation of a device conform to current good manufacturing practice, as prescribed in such regulations, to assure that a device will be safe and effective and otherwise in compliance with federal law.

67. The regulations requiring conformance to good manufacturing practices are set forth in 21 C.F.R. § 820, *et seq.* As explained in the Federal Register, because the Current Good Manufacturing Practice (CGMP) regulations must apply to a variety of medical devices, the regulations do not prescribe the details for how a manufacturer must produce a device. Rather, the quality system regulations provide a framework of basic requirements for each manufacturer to use in establishing a quality system appropriate to the devices designed and manufactured and the

manufacturing processes employed. Manufacturers must adopt current and effective methods and procedures for each device they design and manufacture to comply with and implement the basic requirements set forth in the quality system regulations.

68. Pursuant to 21 C.F.R. § 820.1(c), the failure to comply with any applicable provision in Part 820 renders a device adulterated under section 501(h) of the Federal Drug & Cosmetic Act (“the Act”). 21 U.S.C. § 351.

69. Pursuant to 21 C.F.R. § 820.5, each manufacturer shall establish and maintain a quality system that is appropriate for the specific medical device designed or manufactured. “Quality system” means the organizational structure, responsibilities, procedures, processes and resources for implementing quality management. 21 C.F.R. § 820.3(v).

70. Pursuant to 21 C.F.R. § 820.22, each manufacturer shall establish procedures for quality audits and conduct such audits to assure that the quality system is in compliance with the established quality system requirements and to determine the effectiveness of the quality system.

71. Pursuant to 21 C.F.R. § 820.30(a), each manufacturer shall establish and maintain procedures to control the design of the device in order to ensure that specified design requirements are met.

72. Pursuant to 21 C.F.R. § 820.30(d), each manufacturer shall establish and maintain procedures for defining and documenting design output in terms that allow an adequate evaluation of conformance to design input requirements.

73. Pursuant to 21 C.F.R. § 820.30(e), each manufacturer shall establish and maintain procedures to ensure that formal documented reviews of the design results are planned and conducted at appropriate stages of the device’s design development.

74. Pursuant to 21 C.F.R. § 820.30(f), each manufacturer shall establish and maintain procedures for verifying the device design to confirm that the device design output meets the

design input requirements.

75. Pursuant to 21 C.F.R. § 820.30(g), each manufacturer shall establish and maintain procedures for validating the device design. Design validation shall be performed under defined operating conditions on initial production units, lots or batches, or their equivalents. Design validations shall ensure that devices conform to defined user needs and intended uses and shall include testing of production units under actual or simulated use conditions.

76. Pursuant to 21 C.F.R. § 820.30(h), each manufacturer shall establish and maintain procedures to ensure that the device design is correctly translated into production specifications.

77. Pursuant to 21 C.F.R. § 820.30(i), each manufacturer shall establish and maintain procedures for the identification, documentation, validation or where appropriate verification, review and approval of design changes before their implementation.

78. Pursuant to 21 C.F.R. § 820.70(a), each manufacturer shall develop, conduct, control and monitor production processes to ensure that a device conforms to its specifications. Where deviations from device specifications could occur as a result of the manufacturing process, the manufacturer shall establish and maintain process control procedures that describe any process controls necessary to ensure conformance to specifications. Such process controls shall include:

- (a) documented instructions, standard operating procedures (SOPs) and methods that define and control the manner of production;
- (b) monitoring and control of process parameters and component and device characteristics during production;
- (c) compliance with specified reference standards or codes;
- (d) the approval of processes and process equipment; and
- (e) criteria for workmanship which shall be expressed in documented standards or by means of identified and approved representative samples.

79. Pursuant to 21 C.F.R. § 820.70(b), each manufacturer shall establish and maintain procedures for changes to a specification, method, process or procedure.

80. Pursuant to 21 C.F.R. § 820.70(c), each manufacturer shall establish and maintain procedures to adequately control environmental conditions that could reasonably be expected to have an adverse effect on product quality, including periodic inspection of environmental control system(s) to verify that the system, including necessary equipment, is adequate and functioning properly.

81. Pursuant to 21 C.F.R. § 820.70(e), each manufacturer shall establish and maintain procedures to prevent contamination of equipment or product by substances that could reasonably be expected to have an adverse effect on produce quality.

82. Pursuant to 21 C.F.R. § 820.70(g), each manufacturer shall ensure that all equipment used in the manufacturing process meets specified requirement and is appropriately designed, constructed, placed and installed to facilitate maintenance, adjustment, cleaning and use.

83. Pursuant to 21 C.F.R. § 820.70(h), each manufacturer shall establish and maintain procedures for the use and removal of manufacturing material which could reasonably be expected to have an adverse effect on product quality to ensure that it is removed or limited to an amount that does not adversely affect the device's quality.

84. Pursuant to 21 C.F.R. § 820.70(i), when computers or automated data processing systems are used as part of production or the quality system, the manufacturer shall validate computer software for their intended use according to an established protocol.

85. Pursuant to 21 C.F.R. § 820.72, each manufacturer shall ensure that all inspection, measuring and test equipment, including mechanical, automated or electronic inspection and test equipment, is suitable for the intended purposes and is capable of producing valid results. Each manufacturer shall establish and maintain procedures to ensure that equipment is routinely

calibrated, inspected, checked and maintained.

86. Pursuant to 21 C.F.R. § 820.75(a), where the results of a process cannot be fully verified by subsequent inspection and test, the process shall be validated with a high degree of assurance and approved according to established procedures. “Process validation” means establishing by objective evidence that a process consistently produces a result or product meeting its predetermined specifications. *See* 21 C.F.R. § 820.3(z)(1).

87. Pursuant to 21 C.F.R. § 820.75(b), each manufacturer shall establish and maintain procedures for monitoring and control of process parameters for validated processes to ensure that the specified requirements continue to be met. Each manufacturer shall ensure that validated processes are performed by qualified individuals.

88. Pursuant to 21 C.F.R. § 820.90, each manufacturer shall establish and maintain procedures to control product that does not conform to specified requirements.

89. Pursuant to 21 C.F.R. § 820.100, each manufacturer shall establish and maintain procedures for implementing corrective and preventive action. The procedures shall include requirements for:

- (a) analyzing processes, work operations, concessions, quality audit reports, quality records, service records, complaints, returned product, and other sources of quality data to identify existing and potential causes of nonconforming product or other quality problems;
- (b) investigating the cause of nonconformities relating to product, processes and the quality system;
- (c) identifying the action(s) needed to correct and prevent recurrence of nonconforming product and other quality problems;
- (d) verifying or validating the corrective and preventative action to ensure that

such action is effective and does not adversely affect the finished device;

- (e) implementing and recording changes in methods and procedures needed to correct and prevent identified quality problems;
- (f) ensuring that information related to quality problems or nonconforming product is disseminated to those directly responsible for assuring the quality of such product or the prevention of such problems; and

- (g) submitting relevant information on identified quality problems, as well as corrective and preventative actions, for management review.

90. Upon information and belief, Defendants' One System and DreamStation CPAP devices are adulterated pursuant to 21 U.S.C. § 351 because, among other things, it failed to meet established performance standards and/or the methods, facilities or controls used for its manufacture, packing, storage or installation are not in conformity with federal requirements. *See* 21 U.S.C. § 351.

91. Upon information and belief, Defendants' One System and DreamStation CPAP devices are misbranded because, among other things, it is dangerous to health when used in the manner prescribed, recommended or suggested in the labeling thereof. *See* 21 U.S.C. § 352.

92. Upon information and belief, Defendants' One System and DreamStation CPAP devices are adulterated pursuant to 21 U.S.C. § 351 because Philips failed to establish and maintain CGMP for its One System and DreamStation CPAP devices in accordance with 21 C.F.R. § 820, *et seq.*, as set forth above.

93. Upon information and belief, Philips failed to establish and maintain CGMP with respect to the quality audits, quality testing and process validation for the recalled devices, including the Philips One System and DreamStation CPAP devices.

94. As a result of Philips' failure to establish and maintain CGMP as set forth above, Philips' One System and DreamStation CPAP devices were defective, resulting in injuries to Decedent Robert Kaminsky.

95. If Philips had complied with the federal requirements regarding CGMP, Philips' One System and DreamStation CPAP devices would have been manufactured and/or designed properly such that it would not have resulted in injuries to Decedent Robert Kaminsky.

CAUSES OF ACTION
COUNT I
STRICT PRODUCTS LIABILITY
DEFECTIVE DESIGN

96. Plaintiff incorporates by reference, as if fully set forth herein, each and every allegation set forth in the preceding paragraphs and further alleges as follows:

97. At all times herein mentioned, Defendants designed, researched, manufactured, tested, advertised, promoted, marketed, sold, and/or distributed the Philips DreamStation CPAP devices as hereinabove described that were prescribed to and used by Decedent.

98. Defendants each had a duty to place into the stream of commerce, manufacture, distribute, market, promote and sell the Philips DreamStation CPAP devices so that they were neither defective nor unreasonably dangerous when used for which they were designed, manufactured, distributed, marketed, and sold.

99. At all times herein mentioned, the Philips DreamStation CPAP devices were in an unsafe, defective, and inherently dangerous condition for users such as Plaintiff.

100. At all times of use of the Devices by Decedent, were being used for the purposes and in a manner normally intended, namely for use as treatment for sleep apnea.

101. At the time the Devices left the possession of Defendants and the time the Philips DreamStation CPAP devices entered the stream of commerce, they were in an unreasonably dangerous or defective condition. These defects include, but are not limited to, the following:

- (a) the Devices were not reasonably safe as intended to be used;
- (b) the Devices had an inadequate design for the purpose of treatment of sleep apnea, in that the sound abatement foam should not release toxic and carcinogenic particles and should not have been placed in the devices'

airpath where such particles would then travel directly into patients' lungs and bodies;

- (c) the Devices contained unreasonably dangerous design defects, including an inherently defective design, i.e., placement of a sound abatement foam that releases toxic and carcinogenic particles directly in the airpath of the Devices, from where such particles could easily travel to the user;
- (d) the Devices' defective design resulted in a CPAP device which had risks that far exceeded the benefits of their medical utility;
- (e) the Devices were not appropriately or adequately tested before their distribution; and
- (f) the Devices have an unreasonably high propensity for the release of toxic and carcinogenic particles under normal and expected use of the Devices.
- (g) the Devices have built-in settings for heat and humidity that are expected to be utilized during normal use, and according to Philips such environmental factors may exacerbate the release of toxic and carcinogenic particles from the sound abatement foam in the Devices.

102. At all times herein mentioned, the Devices were expected to and did reach the usual consumers, handlers, and persons coming into contact with said Devices without substantial change in the condition in which they were designed, produced, manufactured, sold, distributed, and marketed by Defendants.

103. The Philips DreamStation CPAP devices' unsafe, defective, and inherently dangerous conditions were the cause of injuries and the ultimate death of Decedent.

104. The Devices failed to perform as safely as an ordinary consumer would expect when used in an intended or reasonably foreseeable manner.

105. Decedent's injuries and death resulted from use of the Devices that were both intended and reasonably foreseeable by Defendants.

106. At the time of Defendants' initial design, manufacture, marketing and sale of the Devices, a feasible, alternative safer design for the Devices were known and available to Philips.

107. At the time of and subsequent to Defendants' initial design, manufacture, marketing and sale of the Devices, including prior to the time of Decedent Robert Kaminsky's initial purchase and use of the Device, Defendants had the ability to eliminate the unsafe character of the Devices without impairing their usefulness, as by either using non-toxic, non-carcinogenic sound abatement foam, or by simply placing the sound abatement foam anywhere else in the Devices besides the Devices' airpath, among other reasonable alternatives.

108. Had Defendants properly and adequately tested the Devices, Defendants would have discovered that the sound abatement foam had a high propensity for releasing toxic and carcinogenic particles when used normally by patients.

109. The Philips DreamStation CPAP devices, manufactured and supplied by Defendants, were, therefore, defective in design or formulation in that, when they left the hands of Defendants, the foreseeable risk of harm from the product exceeded or outweighed the benefit or utility of the Devices' particular design or formulation, and/or it were unreasonably dangerous to the user or consumer, and/or it failed to comply with federal requirements for these medical devices.

110. The foreseeable risks associated with the design or formulation of the Philips DreamStation CPAP devices include, but are not limited to, the fact that the design or formulation of these devices are more dangerous than a reasonably prudent consumer would expect when used

in an intended or reasonably foreseeable manner, and/or it failed to comply with federal requirements.

111. At all times herein mentioned, the Defendants knew, or should have known, that the Devices were in a defective condition, and were inherently dangerous and unsafe for use.

112. Defendants designed, researched, manufactured, tested, advertised, promoted, marketed, sold and distributed defective products which, when used in their intended or reasonably foreseeable manner, created an unreasonable risk to the health of consumers and to Decedent, in particular, and Defendants are therefore strictly liable for the injuries sustained by Decedent.

113. As a direct and proximate result of Decedent Robert Kaminsky's use of Defendants' DreamStation CPAP devices, as manufactured, designed, sold, supplied, marketed and introduced into the stream of commerce by Defendants and/or their failure to comply with federal requirements, Decedent Robert Kaminsky suffered serious physical injury, harm, damages and economic loss, and death. Plaintiff Doris Kaminsky is entitled to compensatory damages in an amount to be determined by the trier of fact.

WHEREFORE, Plaintiff demands judgment against Defendants for compensatory and punitive damages, together with interest, costs of suit, attorneys' fees and all such other relief as the Court deems proper.

COUNT II
STRICT PRODUCTS LIABILITY
FAILURE TO WARN

114. Plaintiff incorporates by reference, as if fully set forth herein, each and every allegation set forth in the preceding paragraphs and further alleges as follows:

115. At all times relevant herein, Defendants were engaged in the design, development, testing, manufacturing, marketing and sale of the Philips DreamStation CPAP devices.

116. Defendants designed, manufactured, assembled and sold the Philips CPAP devices to medical distributors and patients knowing that they would then be used by patients to treat sleep apnea.

117. The Devices placed into the stream of commerce by Defendants were defective due to inadequate warnings because Defendants knew or should have known that the Devices could release toxic and/or carcinogenic particles in patients when used and therefore gives rise to serious physical injury, pain and suffering, debilitation, and death, but failed to give consumers adequate warning of such risks.

118. Defendants had a duty to warn their sales representatives/distributors, prescribing sleep doctors, and patients such as Decedent Robert Kaminsky, and Defendants breached their duty in that they failed to provide adequate and timely warnings or instructions regarding their Philips DreamStation CPAP devices, and their known defects and potential risks, including their propensity to release toxic and/or carcinogenic particles when used normally.

119. Adequate efforts to communicate an adequate warning to the ultimate users were not made by Defendants (or Defendants' sales representatives/distributors).

120. Defendants are strictly liable to Plaintiff because the warnings to Decedent Robert Kaminsky, his medical equipment supplier and his prescribing physician about the dangers the Philips DreamStation CPAP devices posed to consumers when used were inadequate. Examples of the lack and/or inadequacy of Defendants' warnings include, but are not limited to, one or more of the following particulars:

- (a) the Devices contained warnings insufficient to alert Decedent Robert Kaminsky, the medical equipment supplier and Decedent's physicians as to the risk of adverse events, i.e., respiratory issues, development of disease like cancer, and even death, associated with use of the Philips DreamStation CPAP devices, subjecting the Decedent to risks which exceeded the benefits of the Devices;
- (b) the Devices contained warnings insufficient to alert Decedent Robert Kaminsky and his physicians as to the release of toxic and carcinogenic particles when used normally;
- (c) the Devices contained misleading warnings emphasizing the efficacy of the Devices while downplaying the risks associated with their use, thereby making use more dangerous than the ordinary consumer would expect;
- (d) the Devices contained insufficient and/or incorrect warnings to alert consumers, including Decedent Robert Kaminsky, the medical supplier, and the prescribing physicians, regarding the risk, scope, propensity, frequency, duration and severity of the adverse events associated with use of Devices;
- (e) the Devices did not disclose that they were inadequately tested;
- (f) the Devices failed to convey adequate post-marketing warnings regarding the risk, severity, propensity, frequency, scope and/or duration of the dangers posed by normal use of the Devices to treat sleep apnea;
- (g) the Devices failed to contain instructions sufficient to alert consumers to the dangers they posed and to give them the information necessary to avoid or mitigate those dangers.

121. Further, Philips DreamStation CPAP devices are unreasonably dangerous because they were sold to Decedent without an adequate warning that when used normally, the PE-PUR sound abatement foam will release toxic and carcinogenic particles that can lead to serious injury or death.

122. There are other manufacturers of sleep apnea machines on the market that do not contain this foam design defect and Decedent could have chosen to acquire a different model and brand had this defect been disclosed.

123. The Devices placed into the stream of commerce by Defendants were used by patients like Decedent in a manner reasonably anticipated by Defendants.

124. As a direct and proximate result of Defendants' failure to adequately communicate a warning and/or failure to provide an adequate warning and other wrongful conduct as set forth herein, Decedent Robert Kaminsky suffered serious physical injury, harm, damages and economic loss, and death. Plaintiff Doris Kaminsky is entitled to compensatory damages in an amount to be determined by the trier of fact.

WHEREFORE, Plaintiff demands judgment against Defendants for compensatory and punitive damages, together with interest, costs of suit, attorneys' fees and all such other relief as the Court deems proper.

COUNT III
STRICT PRODUCTS LIABILITY
MANUFACTURING DEFECT AND
FAILURE TO ADHERE TO QUALITY CONTROLS

125. Plaintiff incorporates by reference, as if fully set forth herein, each and every allegation set forth in the preceding paragraphs and further allege as follows:

126. The recalled devices, including Decedent's Devices, are defectively manufactured because the foreseeable risks of cancer and other serious injury and illness outweigh the benefits associated with the Devices.

127. The Philips One System and DreamStation CPAP devices were designed and/or manufactured in a manner violative of the Federal Food, Drug and Cosmetic Act, 21 U.S.C. § 321 et seq., and the Medical Devices Amendment thereto (hereafter "FDCA"). The facilities or controls used by defendants in the manufacture, testing, packing, storage, or installation of the Devices were not in conformity with applicable requirements of the FDCA.

128. The Philips One System and DreamStation CPAP devices were expected to and did reach the Decedent without substantial change or adjustment to its function.

129. Defendants knew or should have known of the manufacturing defects and the risk of serious bodily injury that exceeded the benefits associated with the Philips One System and DreamStation CPAP devices.

130. Furthermore, the Philips One System and DreamStation CPAP devices and their defects presented an unreasonably dangerous risk beyond what the ordinary consumer would reasonably expect.

131. The Philips One System and DreamStation CPAP devices were inherently dangerous for the intended use due to a manufacturing defect or defects and improper functioning. Defendants are therefore strictly liable to the Plaintiff for their breach of duty to the Decedent.

132. As a direct and proximate result of Defendants' wrongful conduct, the Decedent sustained severe physical injuries, harm, damages and economic loss, and death. Plaintiff Doris Kaminsky has suffered and will continue to suffer severe emotional distress, mental anguish, economic losses, and other damages for which she is entitled to compensatory damages in an amount to be proven at trial.

WHEREFORE, Plaintiff demands judgment against Defendants for compensatory and punitive damages, together with interest, costs of suit, attorneys' fees and all such other relief as the Court deems proper.

COUNT IV
NEGLIGENCE

133. Plaintiff incorporates by reference, as if fully set forth herein, each and every allegation set forth in the preceding paragraphs and further allege as follows:

134. While the focus of Plaintiff's strict liability claims (Claims I-III) is on the condition of the product, the focus of Plaintiff's negligence claim is instead on Defendants' conduct. Defendants had a duty to exercise reasonable care in the design, formulation, manufacture, testing, quality assurance, quality control, labeling, warning, sale and/or distribution of the Philips DreamStation CPAP devices, including a duty to assure that their products did not pose a significantly increased risk of life-threatening bodily harm and disease.

135. Defendants failed to exercise reasonable care in the design, formulation, manufacture, sale, testing, quality assurance, quality control, labeling, warning, marketing, promotions and distribution of the Philips One System and DreamStation CPAP devices in that Defendants knew or should have known that these products caused significant bodily harm and were not safe for use by consumers.

136. The negligence of Defendants, their agents, servants, and/or employees, included but was not limited to the following acts and/or omissions:

- (a) Negligently designing the recalled devices' PE-PUR sound abatement foam such that it has a high propensity to release toxic and carcinogenic particles during normal use of the Devices;

- (b) Negligently designing the recalled devices such that the sound abatement foam is placed in the airpath of the devices, where the foam's propensity to release toxic and carcinogenic particles is most deleterious to a patient's health because they will directly inhale such toxins and carcinogens;
- (c) Negligently designing the recalled Devices such that they contain built-in settings for use that allow a user to increase the heat and humidity of the air being convected through the devices' airpaths, despite Defendants knowing that heat and humidity can exacerbate the release of the toxic and carcinogenic particles from the PE-PUR sound abatement foam;
- (d) Designing, manufacturing, producing, creating, and/or promoting the Devices for use in treating sleep apnea without adequately, sufficiently, or thoroughly testing them, including both pre-market testing and post-market surveillance;
- (e) Not conducting sufficient testing programs to determine whether or not the PE-PUR sound abatement foam was safe for use in the Devices;
- (f) Selling the devices without making proper and sufficient tests to determine the dangers when used in a reasonably foreseeable and normal manner;
- (g) Negligently failing to adequately and correctly warn Decedent or Decedent's physicians, hospitals, healthcare providers, and medical device distributors of the dangers of using the recalled devices, including:
 - 1) Negligently failing to warn of an increased risk of release of toxic and carcinogenic particles;

- 2) Negligently failing to warn of the risk of development of serious disease such as cancer or even death;
 - 3) Negligently failing to recall their dangerous and defective CPAP devices at the earliest date it became known that the devices were, in fact, dangerous and defective;
 - 4) Negligently advertising and recommending the use of the Devices despite the fact Defendants knew or should have known of their dangerous propensities;
 - 5) Negligently representing that the Devices were safe for their intended use, when in fact, they were unsafe;
 - 6) Negligently manufacturing the Devices in a manner which was dangerous to those individuals who used them;
- (h) Defendants under-reported, underestimated, and downplayed the serious dangers associated with the PE-PUR sound abatement foam used in all of the recalled Devices;
 - (i) Defendants failed to use due care in designing and manufacturing the Devices so as to ensure good performance and durability and reduce the risk of degradation and off-gassing of toxic and carcinogenic particles that could be directly inhaled by the patient;
 - (j) Failed to accompany the Devices with proper warnings;
 - (k) Failed to accompany the Devices with proper instructions for use;

- (l) Failed to conduct adequate testing, including pre-clinical and clinical testing and post-marketing surveillance to determine the safety of the recalled Devices when used normally;
- (m) Were otherwise careless and/or negligent.

137. Despite the fact that Defendants knew or should have known that use of the Philips One System and DreamStation CPAP devices caused harm to individuals that used them, Defendants continued to market, manufacture, distribute and/or sell the Philips One System and DreamStation CPAP devices for use in treating sleep apnea.

138. Defendants knew or should have known that consumers such as Decedent would suffer foreseeable injury, and/or be at increased risk of suffering injury as a result of Defendants' failure to exercise ordinary care, as set forth above.

139. Defendants, furthermore, in advertising, marketing, promoting, packaging and selling the Devices negligently misrepresented material facts regarding their safety, efficacy and fitness for human use by claiming the Devices were fit for their intended purpose of use when, in fact, they were not.

140. Defendants' negligence was the proximate cause of Decedent's physical, mental and emotional injuries, and ultimate death, as well as the economic loss which Plaintiff has suffered and/or will continue to suffer.

141. By reason of the foregoing, Plaintiff experienced and will continue to experience severe harmful effects as a result of the Defendants' negligence as set forth above.

142. Further, as a result of the foregoing acts and omissions, Decedent suffered lost wages.

143. Defendants' conduct, as described above, including, but not limited to, Defendants' failure to adequately test and warn, as well as their continued marketing and distribution of the Philips One System and DreamStation CPAP devices when they knew or should have known of the serious health risks these devices created when used normally by patients such as Decedent.

144. As a direct and proximate result of Defendants' negligence, including negligent testing, failure to warn and misrepresentations, Decedent Robert Kaminsky suffered serious physical injury, harm, damages and economic loss, and ultimately caused his death. Plaintiff Doris Kaminsky will continue to suffer damages and economic loss in the future for which she is entitled to compensatory damages in an amount to be proven at trial.

WHEREFORE, Plaintiff demands judgment against Defendants for compensatory and punitive damages, together with interest, costs of suit, attorneys' fees and all such other relief as the Court deems proper.

COUNT V
NEGLIGENT MISREPRESENTATION

145. Plaintiff incorporates by reference, as if fully set forth herein, each and every allegation set forth in the preceding paragraphs and further allege as follows:

146. Defendants supplied false information to the public, to Decedent and to Decedent's physicians regarding the high-quality, safety and effectiveness of the Philips One System and DreamStation CPAP devices. Defendants provided this false information to induce the public, Decedent and Decedent's physicians to purchase and use the Philips One System and DreamStation CPAP devices.

147. Defendants knew or should have known that the information they supplied regarding the purported high-quality, safety and effectiveness of the Devices would induce

Decedent and Decedent's physicians to purchase and use the Philips One System and DreamStation CPAP devices were false and misleading.

148. Defendants were negligent in obtaining or communicating false information regarding the purported high-quality, safety and effectiveness of the Philips One System and DreamStation CPAP devices.

149. Decedent and Decedent's physicians relied on the false information supplied by Defendants to Decedent's detriment by causing the Philips One System and DreamStation CPAP devices to be purchased and used by Decedent.

150. Decedent and Decedent's physicians were justified in their reliance on the false information supplied by Defendants regarding the purported high-quality, safety and effectiveness of the Philips One System and DreamStation CPAP devices.

151. As a direct and proximate result of Defendants' negligent misrepresentations, Decedent experienced significant damages, including but not limited to physical injury, economic loss, and pain and suffering, and ultimate death caused by the Philips One System and DreamStation CPAP devices.

WHEREFORE, Plaintiff demands judgment against Defendants for compensatory and punitive damages, together with interest, costs of suit, attorneys' fees and all such other relief as the Court deems proper.

COUNT VI
BREACH OF EXPRESS WARRANTY

152. Plaintiff incorporates by reference, as if fully set forth herein, each and every allegation set forth in the preceding paragraphs and further allege as follows:

153. Defendants expressly warranted that the Philips One System and DreamStation CPAP devices were safe and effective medical devices to be used for patients suffering from sleep

apnea.

154. At the time Defendants marketed, sold and/or distributed the Philips One System and DreamStation CPAP devices, they knew that the Devices were intended for human use, and that Decedent Robert Kaminsky was a foreseeable user of the Devices.

155. The express warranties represented by Defendants were a part of the basis for Decedent's use of the Devices, and he and his physician relied on these warranties in deciding to prescribe the Devices.

156. At the time of the making of the express warranties, Defendants had knowledge of the purpose for which the Devices were to be used, and warranted the same to be in all respects safe, effective and proper for such purpose.

157. The Devices do not conform to these express representations as shown by the development of fatal Hodgkin's lymphoma cancer in Decedent Robert Kaminsky.

158. At the time Defendants marketed, sold and/or distributed the recalled Devices, Defendants expressly warranted that the recalled devices were safe for their intended use.

159. Decedent Robert Kaminsky and his prescribing physician reasonably relied upon Defendants' express warranties.

160. Decedent Robert Kaminsky used the Devices for their intended purpose, and in a reasonably foreseeable manner.

161. The Philips One System and DreamStation CPAP devices manufactured and sold by Defendants did not conform to Defendants' express representations because the Devices caused serious injury and death to Decedent Robert Kaminsky when used as recommended and directed. As a direct and proximate result of Defendants' breach of express warranty, Decedent Robert Kaminsky suffered serious physical injury, harm, damages and economic loss, and death. Plaintiff Doris Kaminsky will continue to suffer damages and economic loss in the future and is entitled to

compensatory damages in an amount to be proven at trial.

WHEREFORE, Plaintiff demands judgment against Defendants for compensatory and punitive damages, together with interest, costs of suit, attorneys' fees and all such other relief as the Court deems proper.

COUNT VII
BREACH OF IMPLIED WARRANTIES OF
MERCHANTABILITY AND FOR A PARTICULAR PURPOSE

162. Plaintiff incorporates by reference, as if fully set forth herein, each and every allegation set forth in the preceding paragraphs and further allege as follows:

163. At the time Defendants designed, manufactured, marketed, sold and distributed the Philips One System and DreamStation CPAP devices for use by Decedent Robert Kaminsky, Defendants knew of the use for which these Devices were intended and impliedly warranted these products to be of merchantable quality and safe for such use and that their design, manufacture, labeling and marketing complied with all applicable federal requirements.

164. The Philips One System and DreamStation CPAP devices manufactured and supplied by Defendants were not of merchantable quality and were not fit for the ordinary and/or particular purpose for which they were intended as, among other defects, the risks included an unreasonably high risk of developing cancer or other serious illness due to the release of toxic and carcinogenic particles from the Devices' PE-PUR sound abatement foam.

165. Decedent Robert Kaminsky and/or his physician reasonably relied upon the skill and judgment of Defendants as to whether the Philips One System and DreamStation CPAP devices were of merchantable quality and safe for their intended and particular use and purpose, and upon Defendants' implied warranty as to such matters.

166. Contrary to such implied warranties, the Philips One System and DreamStation CPAP devices were not of merchantable quality or safe for their intended and particular use and purpose, because the Devices were defective when used normally as described above, and/or failed to comply with federal requirements.

167. As a direct and proximate result of Defendants' breach of implied warranties, Decedent Robert Kaminsky suffered serious physical injury, harm, damages, economic loss and ultimately death. Plaintiff Doris Kaminsky will continue to suffer harm, damages and economic loss in the future and is entitled to compensatory damages in an amount to be proven at trial.

WHEREFORE, Plaintiff demands judgment against Defendants for compensatory and punitive damages, together with interest, costs of suit, attorneys' fees and all such other relief as the Court deems proper.

COUNT VIII
BREACH OF IMPLIED WARRANTY OF FITNESS FOR A
PARTICULAR PURPOSE

168. Plaintiff incorporates by reference, as if fully set forth herein, each and every allegation set forth in the preceding paragraphs and further alleges as follows:

169. Defendants designed, manufactured, tested, marketed and distributed into the stream of commerce the Philips One System and DreamStation CPAP devices.

170. At the time Defendants designed, manufactured, tested, marketed and distributed into the stream of commerce the Philips One System and DreamStation CPAP devices, Defendants knew the use for which the Philips One System and DreamStation CPAP devices were intended, and impliedly warranted the Philips One System and DreamStation CPAP devices to be safe for such use.

171. Decedent Robert Kaminsky and/or his physician reasonably relied upon the skill and judgment of Defendants as to whether the Philips One System and DreamStation CPAP devices were safe for their intended use.

172. Contrary to Defendants' implied warranties, the Philips One System and DreamStation CPAP devices were not fit for their intended and particular use and purpose, because the Devices were defective when used as described above, and/or failed to comply with federal requirements.

173. As a direct and proximate result of Defendants' breach of implied warranties, Decedent Robert Kaminsky suffered serious physical injury, harm, damages, economic loss and ultimately death. Plaintiff Doris Kaminsky will continue to suffer harm, damages and economic loss in the future and is entitled to compensatory damages in an amount to be proven at trial.

WHEREFORE, Plaintiff demands judgment against Defendants for compensatory and punitive damages, together with interest, costs of suit, attorneys' fees and all such other relief as the Court deems proper.

COUNT IX
VIOLATION OF THE NEW YORK DECEPTIVE TRADE PRACTICES ACT
(N.Y. GEN. BUS. LAW §§ 349 ET SEQ.; 350-e ET SEQ.)

174. Plaintiff incorporates by reference, as if fully set forth herein, each and every allegation set forth in the preceding paragraphs and further alleges as follows:

175. Defendants unfairly, unconscionably, and deceptively advertised, marketed, sold, and represented the Philips One System and DreamStation CPAP devices as a high-quality, safe and effective medical devices for treatment of sleep apnea to Decedent and Decedent's physicians.

176. Before they advertised, marketed, sold and represented the Philips One System and DreamStation CPAP devices that were used by Decedent, Defendants knew or should have known of the unreasonable dangers and serious health risks that such Devices posed to patients like

Decedent.

177. Decedent purchased and used the Philips One System and DreamStation CPAP devices for personal use and thereby suffered ascertainable losses as a result of Defendants' actions in violation of the consumer protection laws.

178. Had Defendants not engaged in the deceptive conduct described herein, Decedent would not have purchased and/or paid for the Philips One System and DreamStation CPAP devices, and would not have incurred related medical costs and fatal injuries.

179. Defendants engaged in wrongful conduct while at the same time obtaining, under false pretenses, moneys from Decedent for the Philips One System and DreamStation CPAP devices that would not have been paid had Defendants not engaged in unfair and deceptive conduct.

180. Unfair methods of competition or deceptive acts or practices that are proscribed by law, include the following:

- (a) Representing that goods or services have characteristics, ingredients, uses, benefits or quantities that they do not have;
- (b) Advertising goods or services with the intent not to sell them as advertised; and
- (c) Engaging in fraudulent or deceptive conduct that creates a likelihood of confusion or misunderstanding.

181. Decedent was injured and subsequently killed by the cumulative and indivisible nature of Defendants' conduct. The cumulative effect of Defendants' conduct directed at patients, physicians and consumers was to create demand for and sell the Philips One System and DreamStation CPAP devices. Each aspect of Defendants' conduct combined to artificially create sales of the Philips One System and DreamStation CPAP devices.

182. Defendants have a statutory duty to refrain from unfair or deceptive acts or trade practices in the design, development, manufacture, promotion and sale of the Philips One System and DreamStation CPAP devices.

183. Had Defendants not engaged in the deceptive conduct described above, Decedent would not have purchased and/or paid for the Philips One System and DreamStation CPAP devices, and would not have incurred related medical costs.

184. Defendants' deceptive, unconscionable, or fraudulent representations and material omissions to patients, physicians and consumers, including Decedent, constituted unfair and deceptive acts and trade practices in violation of the state consumer protection statutes listed.

185. Defendants' actions, as complained of herein, constitute unfair competition or unfair, unconscionable, deceptive or fraudulent acts, or trade practices in violation of state consumer protection statutes, as listed below.

186. Defendants have engaged in unfair competition or unfair or deceptive acts or trade practices or have made false representations in violation of the New York Deceptive Trade Practices Act, (N.Y. Gen. Bus. Law §§ 349 *et seq.*; 350-e *et seq.*).

187. Under the statutes listed above to protect consumers against unfair, deceptive, fraudulent and unconscionable trade and business practices and false advertising, Defendants are the suppliers, manufacturers, advertisers, and sellers, who are subject to liability under such legislation for unfair, deceptive, fraudulent and unconscionable consumer sales practices.

188. Defendants violated the statutes that were enacted to protect consumers against unfair, deceptive, fraudulent and unconscionable trade and business practices and false advertising, by knowingly and falsely representing that the Philips One System and DreamStation CPAP devices were fit to be used for the purpose for which they were intended, when in fact these Devices were defective and

dangerous, and by other acts alleged herein. These representations were made in uniform promotional materials.

189. The actions and omissions of Defendants alleged herein are uncured or incurable deceptive acts under the statutes enacted to protect consumers against unfair, deceptive, fraudulent and unconscionable trade and business practices and false advertising.

190. Defendants had actual knowledge of the defective and dangerous condition of the Philips Philips One System and DreamStation devices and failed to take any action to cure such defective and dangerous conditions.

191. Decedent and the medical community relied upon Defendants' misrepresentations and omissions in determining which CPAP / sleep apnea treatment devices to use and recommend.

192. Defendants' deceptive, unconscionable or fraudulent representations and material omissions to patients, physicians and consumers, constituted unfair and deceptive acts and practices.

193. By reason of the unlawful acts engaged in by Defendants, and as a direct and proximate result thereof, Decedent suffered ascertainable losses and damages.

194. As a direct and proximate result of Defendants' violations of New York's consumer protection laws, Plaintiff has sustained economic losses and other damages and is entitled to statutory and compensatory, damages in an amount to be proven at trial.

195. As specifically described in detail above, Defendants knew that the Philips One System and DreamStation CPAP devices subjected patients to the release of toxic and carcinogenic particles leading to serious illness, injury, and even death.

196. As a direct and proximately result of Defendants' representations, Decedent has experienced significant damages, including but not limited to permanent physical injury, economic loss, pain and suffering, the need for sustained medical treatment and observation, and ultimate

death, caused by the Philips DreamStation CPAP devices.

WHEREFORE, Plaintiff demands judgment against Defendants for compensatory and punitive damages, together with interest, costs of suit, attorneys' fees and all such other relief as the Court deems proper.

COUNT X
PUNITIVE DAMAGES

197. Plaintiff incorporates by reference, as if fully set forth herein, each and every allegation set forth in the preceding paragraphs and further alleges as follows:

198. Defendants risked the safety of recipients of their products, including Decedent, with knowledge of the safety and efficacy problems and suppressed this knowledge from the general public.

199. Defendants made conscious decisions not to redesign, re-label, warn or inform the unsuspecting recipients of its recalled Devices despite knowledge that these Devices were defective and unreasonably dangerous in nature.

200. Defendants knew or ought to have known that this conduct would result in injury or death, but continued to mislead both the medical community and the public at large, including Decedent, by making false representations about the safety and efficacy of the recalled devices.

201. These acts are wanton and reckless in that the Defendants demonstrated conscious indifference and utter disregard of the consequences of their actions upon the health, safety and rights of others, including Decedent.

202. Additionally, Defendants delayed the recall of the defective devices while seeking clearance for the next-generation DreamStation 2 device, which is significantly more expensive

than the recalled first-generation devices, and did not disclose to the public any of the risks described herein until after the DreamStation 2 had been made commercially available. Thus, Defendants allowed patients like Decedent to continue to be exposed to toxic and carcinogenic particles for a significantly longer period of time while Defendants were attempting to monetize this public health crisis of their own creation.

203. As a direct and proximate result of Defendants' conscious and deliberate disregard for the rights and safety of consumers such as Decedent, Decedent suffered severe and permanent physical injuries, and ultimately death, as set forth above. Defendants' outrageous conduct warrants an award of punitive damages.

WHEREFORE, Plaintiff demands judgment against defendants for compensatory, treble, and punitive damages, together with interest, costs of suit, attorneys' fees, and all such other relief as the Court deems proper.

PRAYER FOR RELIEF

WHEREFORE, Plaintiff prays for the following relief:

1. Judgment in favor of Plaintiff and against all Defendants, for damages in such amounts as may be proven at trial;
2. Compensation for both economic and non-economic losses, including but not limited to medical expenses, loss of earnings, disfigurement, pain and suffering, mental anguish, emotional distress, and loss of consortium, in such amounts as may be proven at trial;
3. Punitive and/or exemplary damages in such amounts as may be proven at trial;
4. Attorneys' fees and costs;
5. Interest; and
6. Any and all further relief, both legal and equitable, that the Court may deem just and proper.

Dated: June 26, 2023

MERSON LAW, PLLC

/s/ Nathan Werksman

Jordan K. Merson, Esq.
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JURY TRIAL DEMANDED

Plaintiff demands a trial by jury of all issues.

Dated: June 26, 2023

MERSON LAW, PLLC

/s/ Nathan Werksman

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